UGI Development Company

390 Route 11 P.O. Box 224 Hunlock Creek, PA 18621-0224 (570) 830-1269

March 7, 2006

CERTIFIED MAIL 7002 0860 0000 1118 1942

Mrs. Rene McLaughlin 3 AP 2 United States Environmental Protection Agency Region 3 1650 Arch Street Philadelphia, PA 19103-2025



RE: UGI Development Company

RATA Notification - Units No. 4 and 6

Dear Mrs. McLaughlin:

This letter is to notify you that UGI Development Company is tentatively scheduled to perform the Annual RATA Testing on Hunlock Power Station's Units No. 6 (Coal Unit) and No. 4 (Combustion Turbine) for the week of April 3, 2006.

If you should have any questions, please call me at (570) 830-1267.

Sincerely,

Jeffrey T. Steeber Staff Engineer

Hunlock Power Station

JTS:chet



AIR QUALITY TESTING SERVICES

UGI DEVELOPMENT COMPANY HUNLOCK STATION UNIT 4 CT

CEMS RATA PROTOCOL

Catalyst Air Management, Inc. Project Number 124-132

FEBRUARY 16, 2006



UGI DEVELOPMENT COMPANY HUNLOCK STATION UNIT 4 CT

CEMS RATA PROTOCOL

CATALYST AIR MANAGEMENT, INC. PROJECT NUMBER 124-032

February 16, 2006

Prepared for Mr. Jeff Stever UGI Development Company Hunlock Station US Route 11, PO Box 224 Hunlock Creek, PA 18621



STATEMENT OF VALIDITY

UGI Development Company – Hunlock Station Catalyst Project 124-032 February 16, 2006

To the best of our knowledge, the applicable state and federal regulations, operating permits, or plan approvals applicable to each source and control device being tested have been reviewed and all the testing requirements have been incorporated into this test protocol.

Michael J. Taylor

President - Catalyst Air Management

Jeff Steeber

UGI Development Company

All Stockes

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PROJECT FACT SHEET

NAME OF SOURCE OWNER: UGI Development Company

SOURCE IDENTIFICATION: Hunlock Station Unit 4

LOCATION OF SOURCE: Route 11, PO Box 224
Hunlock Creek, PA 18621

TYPE OF OPERATION: Combustion Turbine

TYPES OF TESTS PERFORMED: Oxygen/Carbon Dioxide-EPA Method 3A

Nitrogen Oxide-EPA Method 7E

CEMS DESCRIPTION: TECO 42CHL NOx Sevomex 1440C O₂

Sevollex 1440C O2

TEST COMPANY: Catalyst Air Management, Inc.

2505 Byington-Solway Road Knoxville, TN 37931

SITE SUPERVISOR: Jeff Ferguson - Principal

TEST PERSONNEL: Mark Williams - Technician

PROPOSED TEST DATES: April 3, 2006

OWNERS REPRESENTATIVE: Jeff Steeber

TEST OBSERVER:

1.0 Introduction

Catalyst Air Management, Inc. (Catalyst) has been contracted by UGI Development Company (UGI) to perform the annual Continuous Monitoring System (CEMS) Relative Accuracy Test Audit (RATA) for the CT at the Hunlock Station in Hunlock Creek, PA.

The sampling program is scheduled for the week of April 3, 2006. The Catalyst contact is Mike Taylor (865) 531-0075 or Jeff Ferguson (610) 913-7600. Mr. Jeff Steeber of UGI will coordinate plant and CEMS operation during the testing.

2.0 Description of Process

The Hunlock Station, CT, is a GE LM6000PC natural gas fired turbine operating in the simple cycle mode. The turbine is equipped with water injection in the combustion zone for NOx control. A process diagram is included in the figure section.

3.0 Description of Sampling Locations

The stack is 75 feet high, with an inside diameter of 9.0 feet. The sampling location is approximately 10 feet upstream (1.1 diameters) from the stack exit and 50 feet downstream (5.6 diameters) of the inlet duct. A schematic of the sampling location is included in the figures section.

4.0 Description of CEMS

The CEMS is a heated extractive system that measures NOx and O₂ concentrations at the sampling location. The CEMS analyzers includes a Thermo Environmental Model 42CHL NOx analyzer and a Servomex Model 1440C O₂ analyzer. The recording and reporting requirements are performed by a computerized data acquisition and handing system.

CEMS

- (1) Thermo Environmental 42CHL NOx
- (1) Servomex 1440C O₂

The analyzers measure on a dry basis. The data acquisition and handling system utilizes a Fo factor of 8710 scf/mmBtu to calculate NOx emissions in lbs/mmBtu.

5.0 Sampling Program Procedures

The following test methods were utilized during the test program:

EPA Method 3A Gas Analysis for CO₂, O₂, Excess Air and Dry Molecular Weight

(Instrumental Analyzer Method)

EPA Method 7E Determination of Nitrogen Oxides Emissions from Stationary

Sources (Instrumental Analyzer Method)

5.1 NOx, SO₂ and CO₂ - EPA Methods 3A and 7E

A sample is continuously extracted and introduced into a Thermo Environmental Model 10, Chemiluminescent NOx analyzer and Servomex 1400 O₂/CO₂ analyzer for determination of gas concentrations. The sample is extracted through a heated stainless steel probe, heated sample line and sample conditioner to dry the sample before it enters the analyzers. A sample flow control system is used to control the flow into the analyzers. The analyzers are calibrated prior to starting the testing with EPA Protocol 1, calibration gases. A system bias check is performed before each run by introducing the zero and upscale gas at the back end of the sample probe. The system bias check is repeated at the end of each test run to determine the analyzer zero and calibration drift.

Nine (9) to twelve (12) test runs will be conducted with each being twenty-one (21) minutes in duration. A three (3) point traverse will performed during each run.

The NOx analyzer span as expected to be 0-25 ppm. The O_2/CO_2 analyzer spans are 0-25% and 0-20%, respectively. The calibration gases that will be utilized are zero, 40-60% and 80-100% of span.

Reference	Method	Analyzers:
-----------	--------	------------

Manufacturer	Model	Pollutant	Span
TECO	10A	NOx	0-25 ppm
Servomex	1400B	CO_2/O_2	0-20%/0-25%

The Catalyst reference method values are on a dry basis. The reference method emission rates, in lbs/mmBtu, will be calculated using the measured O₂ values and equation 19-1 in EPA Method 19. The reference method emission rate (lbs/mmBtu) will be compared to the CEMS lbs/mmBtu as recorded by the plant data acquisition system and the relative accuracy calculated.

The CEMS heat input (mmBtu/hr) will be used to determine the NOx emission rate (lb/hr) to compare to the CEMS lb/hr as recorded by the data acquisition system and the relative accuracy calculated.

All the procedures used for the RATA program were performed in accordance with the Code of Federal Regulations, Title 40, Part 60, Appendix A, and Appendix B, Performance Specifications 2, 3 and 6, Part 75 and the Source Testing Manual Rev 3.3.

6.0 Quality Assurance Procedures

The quality assurance procedures followed during the testing activities followed guidelines set forth by the previous mentioned methods and the EPA Quality Assurance Handbook for Source Sampling. The specific procedures for this test program are listed below.

6.1 Instrumental Methods

Analyzer calibrations, system bias check and drift checks were completed before and after each sample run utilizing EPA Protocol 1 calibration gases.

The analyzer interference responses were determined in accordance with Section 5.4 of Method 20 and Section 7.2 through 7.6 of Method 6C.

The NOx analyzer NO₂ to NO converter efficiency is determined in accordance with Section 5.6 of Method 20.

7.0 Unit Operational Data

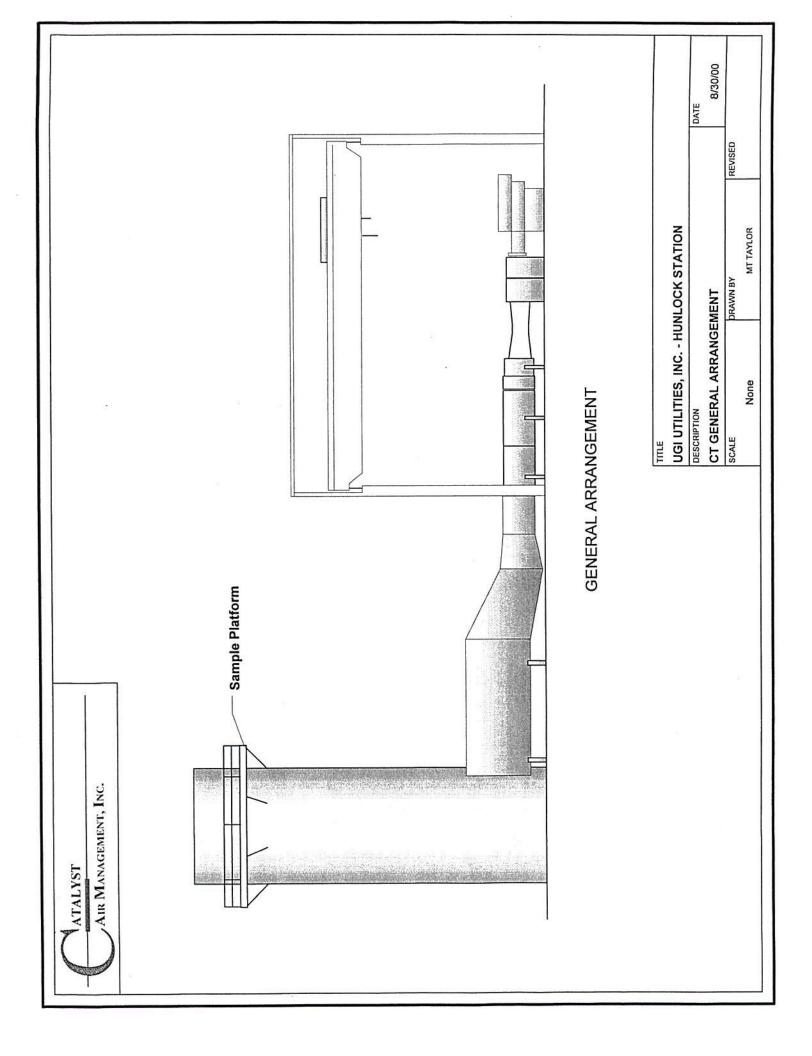
UGI will be responsible for the operation of the CT and CEMS. UGI personnel will provide documentation for operational conditions maintained during the testing. The CEMS data acquisition and handling system (DAHS) will provide emissions data to determine the system relative accuracy. The testing will be performed with the unit operating at 80-100% load.

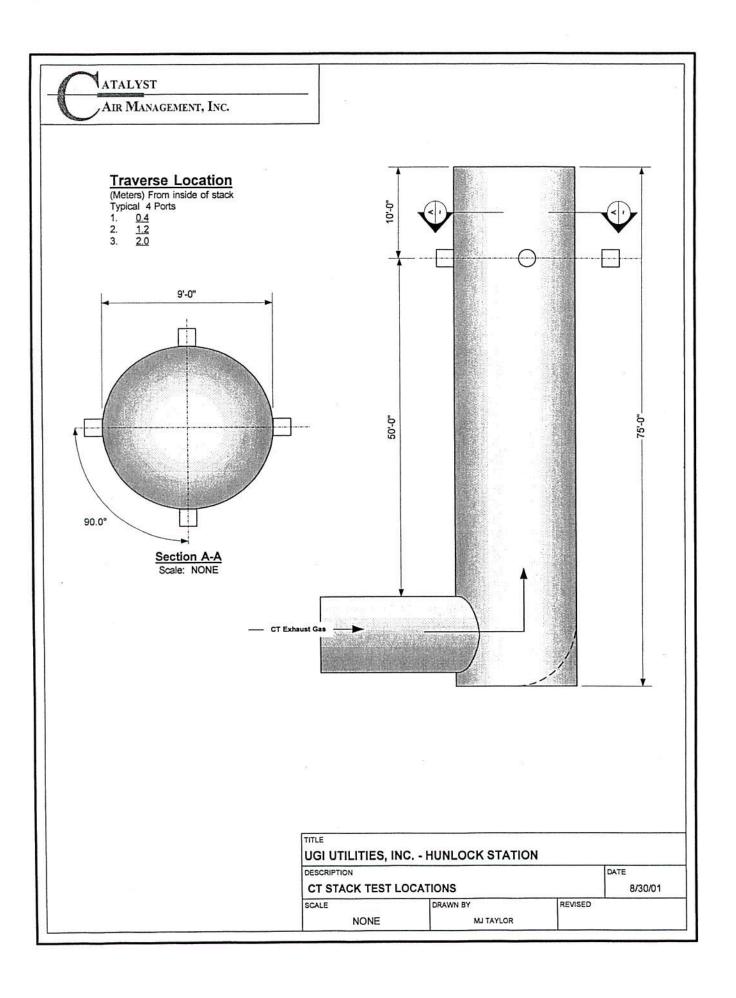
8.0 Test Report

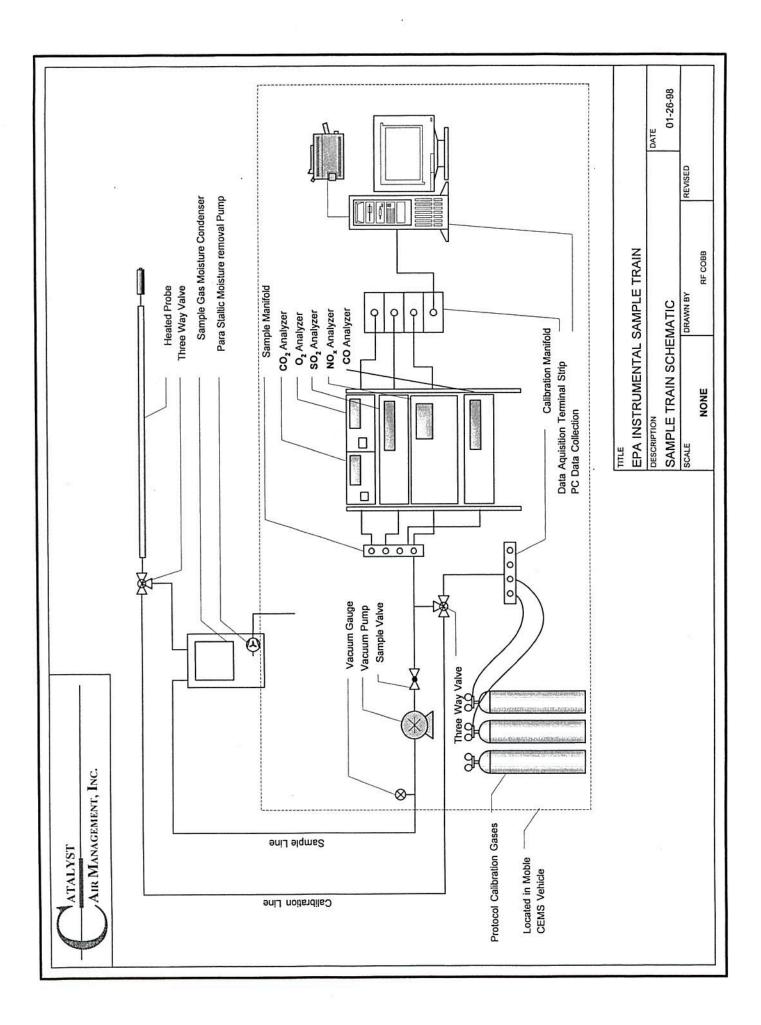
The test report will be submitted that will provide the results of the test program and all the pertinent information pertaining to the individual tests. The report will include the following:

- Unit description
- Sampling location and test points
- Unit operating data and production data
- Test personnel
- Test results and discussion of the test program
- Raw test data
- Equipment calibration and test method QA
- Sample calculations

Figures







Field Data Sheets

CATALYST AIR M. TESTING SERVICE			с.		DA	ГЕ:			RUN	T#:	_
CLIENT:				_	PLA	NT:			UNI	т:	_
Analyzer:	8	_	O2	_			CO2			NOx	
Scale:				_		2400				-	
			7	- =		03	~	Ħ		~~~	_ =
	TIME:	Zero Gas:	Mid-Range:	High-Range:	Zero Gas:		Mid-Range:	High-Range:	Zero Gas:	Mid-Range:	High-Range:
Cylinder Value	L					<u></u>					
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Run 3 Start: End:					_						
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End: End: System Calib. Run 12 Start:										I	
End:[*

Sample Calculations

CALCULATIONS FOR GAS CONCENTRATION

GAS CONCENTRATION (Cgas)

$$C_{gas} = \left(\overline{C} - C_0\right) \left(\frac{C_{ma}}{C_m - C_a}\right)$$

C_{gas} = Effluent gas concentration, ppm

 \overline{C} = Average gas concentration indicated by gas analyzer, dry basis, ppm

C₀ = Average of initial and final system calibration bias check responses for the zero gas, ppm

C_m = Average of initial and final system calibration bias check responses for the upscale calibration gas, ppm

 C_{ma} = Actual concentration of the upscale calibration gas, ppm

CALCULATION FOR EMISSION RATE

$$E(lb/mmbtu) = C \times F_d \left(\frac{20.9}{20.9 - \%O_2} \right)$$

$$E(lb/mmbtu) = C \times F_c(100/CO_2)$$

Where:

C(lb/dscf) = Pollutant concentration (ppm) x conversion factor.

Conversion Factors:

NOx =
$$1.194 \times 10^{-7}$$

SO₂ = 1.660×10^{-7}
CO = 7.274×10^{-8}
C₃H₈ = 1.145×10^{-7}
CH₄ = 4.152×10^{-8}

 $F_d(dscf/mmbtu) = "F"$ Factor for fuel type, (Ref. EPA Method 19)

$$F_d$$
 (Coal) = 9780 F_c (Coal) = 1800 F_d (Gas) = 8710 F_c (Gas) = 1040 F_d (Oil) = 9190 F_c (Oil) = 1420

2. Heat input HI (mmBtu/hr) HI = fuel flow (cf/min) * Heat Value (mmBtu/cf)*60

HI = fuel flow (gal/min) * Heat Value (mmBtu/gal) * 60

3. Emission Rate E(lb/hr) E(lb/hr) = HI (mmBtu/hr) * E(lb/mmBtu)

CALCULATION FOR RELATIVE ACCURACY

ARITHMETIC MEAN (of difference, d, of a data set)

$$d = \frac{1}{n} \sum_{i=1}^{n} d_i$$

Where n =Number of Data points.

ALGEBRAIC SUM (of the individual differences, d_i)

$$\sum_{i=1}^{n} d_{i}$$

STANDARD DEVIATION S_d

$$S_{d} = \left[\frac{\sum_{i=1}^{n} d_{i}^{2} - \left(\sum_{i=1}^{n} d_{i}\right)^{2}}{n} \right]^{\frac{1}{2}}$$

CONFIDENCE COEFFICIENT, CC

$$CC = t_{0.025} \frac{S_d}{\sqrt{n}}$$

For 9 tests $t_{0.975} = 2.306$ For 10 tests $t_{0.975} = 2.262$ For 11 tests $t_{0.975} = 2.228$ For 12 tests $t_{0.975} = 2.201$

RELATIVE ACCURACY, RA

$$RA = \frac{\left| \overline{d} \right| + \left| CC \right|}{\overline{RM}} \times 100$$



AIR QUALITY TESTING SERVICES

UGI DEVELOPMENT COMPANY HUNLOCK STATION UNIT 6

CEMS RATA PROTOCOL

Catalyst Air Management, Inc. Project Number 124-131

FEBRUARY 16, 2006



UGI DEVELOPMENT COMPANY HUNLOCK STATION UNIT 6

CEMS RATA PROTOCOL

CATALYST AIR MANAGEMENT, INC. PROJECT NUMBER 124-131

February 16, 2006

Prepared for Mr. Jeff Steeber UGI Development Comapny Hunlock Station US Route 11, PO Box 224 Hunlock Creek, PA 18621



STATEMENT OF VALIDITY

UGI Development Company – Hunlock Station Catalyst Project 124-031 February 16, 2006

To the best of our knowledge, the applicable state and federal regulations, operating permits, or plan approvals applicable to each source and control device being tested have been reviewed and all the testing requirements have been incorporated into this test protocol.

Michael J. Taylor

President - Catalyst Air Management

Jeff Steeber

UGI Development Company

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PROJECT FACT SHEET

NAME OF SOURCE OWNER:	UGI Development Company
SOURCE IDENTIFICATION:	Hunlock Station Unit 6
LOCATION OF SOURCE:	Route 11, PO Box 224 Hunlock Creek, PA 18621
TYPE OF OPERATION:	Coal Fired Utility Steam Generator
TYPES OF TESTS PERFORMED:	Sample and Velocity Traverse-EPA Method 1 Volumetric Flow Rate-EPA Method 2F/H Oxygen/Carbon Dioxide-EPA Method 3A Moisture Content-EPA Method 4 Sulfur Dioxide-EPA Method 6C Nitrogen Oxide-EPA Method 7E Opacity-EPA Method 9
CEMS DESCRIPTION:	Air Monitor Mastron Flow - B26354 Thermo Environmental SO ₂ - 43C-69110-362 Thermo Environmental NOx - 42C-69062-362 Thermo Environmental CO ₂ - CHL-68209-359 Air Monitor RAC Opacity - 0045-65, 0018-65 VIM Technologies DAHS - 14787WIN
TEST COMPANY:	Catalyst Air Management, Inc. 2505 Byington-Solway Road Knoxville, TN
SITE SUPERVISOR:	Jeff Ferguson - Principal
TEST PERSONNEL:	Rick Derrera - Lead Technician Mark Williams - Technician
PROPOSED TEST DATES:	April 3, 2006
OWNERS REPRESENTATIVE:	Jeff Steeber

TEST OBSERVER:

1.0 Introduction

Catalyst Air Management, Inc. (Catalyst) has been contracted by UGI Development Company (UGI) to perform the Continuous Monitoring System (CEMS) Relative Accuracy Test Audit (RATA) for Unit 6 at the Hunlock Station in Hunlock Creek, PA.

The sampling program is scheduled for the week of April 3, 2006. The Catalyst contact is Mike Taylor (865) 531-0075 or Jeff Ferguson (610) 913-7600. Mr. Jeff Steeber of UGI will coordinate plant and CEMS operation during the testing.

2.0 Description of Process

The Hunlock Station, Unit 6, is a pulverized coal fired boiler. No.2 fuel oil is used for ignition and flame stabilization. The boiler is is rated at 400,000 lbs/hr of steam at 1350 psig and 955 °F. It is a Foster Wheeler drum type, wet bottom, arch fired, balanced draft unit with a nominal heat input of 562 mmBtu/hr. The steam is supplied to a turbine/generator rated at approximately 50 MW. The flue gas is passed through a primary and secondary precipitator for particulate control. A process flow diagram is included in the figures section.

3.0 Description of Sampling Locations

The Unit 6 stack is 40.5 feet high, with an inside diameter of 9.5 feet. The sampling location is approximately 14.5 upstream from the stack exit and 27 feet downstream of the inlet duct. A schematic of the sampling location is included in the figures section.

4.0 Description of CEMS

The Unit 6 CEMS is a dilution extraction system that measures SO₂, NOx and CO₂ concentrations at the sampling location. The CEMS analyzers includes a Thermo Environmental Model 43H SO₂ analyzer, Thermo Environmental Model 42D NOx analyzer, Thermo Environmental Model 41CHL CO₂ analyzer and a Air Monitor Mastron Electron Flow monitor. The recording and reporting requirements are performed by a computerized data acquisition and handing system.

Unit 6 CEMS

- (1) Thermo Environmental SO₂ 43C Serial No. 43C-69110-362
- (1) Thermo Environmental NOx 42D Serial No. 42D-69062-362
- (1) Thermo Environmental CO₂ 41CHL Serial No. 41CHL-68209-359
- (1) Air Monitor Mastron Electronic Flow- Serial No. B26354
- VIM Technologies Data Acquisition and Handling System Serial No.14787WIN

The analyzers measure on a wet basis. The data acquisition and handling system utilizes a Fc factor of 1910 scf/mmBtu to calculate NOx emissions in lbs/mmBtu.

5.0 Sampling Program Procedures

The following test methods were utilized during the test program:

EPA Method 1	Sample and Velocity Traverse for Stationary Sources
EPA Method 2F	Determination of Stack Gas Velocity and Volumetric Flow Rate
	with Three-dimensional Probes
EPA Method 2H	Determination of Stack Gas Velocity and Volumetric Flow Rate
	Taking into Account Velocity Decay Near the Stack Wall
EPA Method 3A	Gas Analysis for CO2, O2, Excess Air and Dry Molecular Weight
	(Instrumental Analyzer Method)
EPA Method 4	Determination of Moisture Content in Stack Gas
EPA Method 6C	Determination of Sulfur Dioxide Emissions from Stationary
	Sources (Instrumental Analyzer Method)
EPA Method 7E	Determination of Nitrogen Oxides Emissions from Stationary
	Sources (Instrumental Analyzer Method)
EPA Method 9	Visual Determination of the Opacity of Emissions from Stationary
	Sources

5.1 NOx, SO₂ and CO₂ - EPA Methods 3A, 6C and 7E

A sample is continuously extracted and introduced into a Thermo Environmental Model 10, Chemiluminescent NOx analyzer, a Western Research Model 721, SO₂ analyzer and Servomex 1400 O₂/CO₂ analyzer for determination of gas concentrations. The sample is extracted through a heated stainless steel probe, heated sample line and sample conditioner to dry the sample before it enters the analyzers. A sample flow control system is used to control the flow into the analyzers. The analyzers are calibrated prior to starting the testing with EPA Protocol 1, calibration gases. A system bias check is performed before each run by introducing the zero and upscale gas at the back end of the sample probe. The system bias check is repeated at the end of each test run to determine the analyzer zero and calibration drift. Nine (9) to twelve (12) test runs will be conducted with each being twenty-one (21) minutes in duration.

The NOx analyzer spans are expected to be 0-250 ppm. The SO_2 analyzer span is expected to be 0-1000. The O_2/CO_2 analyzer spans are 0-25% and 0-20%, respectively. The calibration gases that will be utilized are zero, 40-60% and 80-100% of span.

itelei chec Michiga Analyzers.	Reference	Method	Analyzers:
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Manufacturer	Model	Pollutant	Span
TECO	10A	NOx	0-250 ppm
Western Research	721	SO_2	0-1000 ppm
Servomex	1400B	CO_2/O_2	0-20%/0-25%

5.2 Volumetric Flow Rate - EPA Method 2F/H

Traverse points and flow relative accuracy runs will be performed using EPA Methods 1, 2F, 2H, 3A and 4. The velocity and volumetric flow rate will be determined in accordance with procedures outlined in EPA Method 2F. The wall effect default value of 0.995 will be utilized according to EPA Method 2H. A DAT 3-D probe is connected to a series of calibrated magnehelics and manometers to determine the yaw and pitch angles, static pressure and pitch coefficient of the flue gas at each traverse point. The temperature of the flue gas is also determined at each traverse point.

A minimum of nine (9) test runs will be performed at each load condition. A moisture will be performed for every three (3) flow traverses.

5.3 Moisture - EPA Method 4

The moisture content will be determined in accordance with procedures outlined in EPA Method 4. The flue gas sample is extracted from the gas stream and the moisture content is determined by measuring the increase in volume of the impingers. The sampling train consists of the following equipment connected in series:

Stainless steel heated sample probe

A modified Greenburg-Smith impinger containing 100 ml of H₂O

A Greenburg-Smith impinger containing 100 ml of H₂O

A modified Greenburg-Smith impinger, empty

A modified Greenburg-Smith impinger containing approximately 250g of silica gel

The sample volume is measured by passing it through a calibrated dry gas meter. After the run, the impinger contents are measured for increase in volume. The silica gel is returned to the original tared container and weighed to determine moisture gain.

5.4 F Factor Determination

Coal samples will be collected during the testing in order to determine the dry F factor used to calculate the emission rates in lb/mmBtu.

5.5 Opacity

Catalyst will also conduct an opacity monitor audit by performing nine (9) visible emissions evaluations. The visible emission evaluations will be 15 minutes in duration. Each of the evaluations will generate 15 - one minute averages that will be compared to the one minute averages generated by the CEMS during the same time period. A certified visible emission evaluator will conduct the audit.

All the procedures used for the RATA program were performed in accordance with the Code of Federal Regulations, Title 40, Part 60, Appendix A, and Appendix B, Performance Specifications 2, 3 and 6, Part 75 and the Source Testing Manual Rev 3.3.

6.0 Quality Assurance Procedures

The quality assurance procedures followed during the testing activities followed guidelines set forth by the previous mentioned methods and the EPA Quality Assurance Handbook for Source Sampling. The specific procedures for this test program are listed below.

6.1 Velocity/Volumetric Flow Rate

The 3-D probe will be visually inspected and calibrated to meet the design specifications of EPA Method 2F.

All legs of the probe will be leaked checked before and after each sample run.

The stack thermocouples will be calibrated prior to the testing and a post test check will be performed after the testing project.

The magnehelics will be leveled and zeroed before each sample run.

6.2 Moisture

The dry gas meter is fully calibrated annually using an EPA intermediate standard.

Post -test dry gas meter checks will be completed to verify the accuracy of the meter Yi.

Pre-test and post-test leak checks will be completed and were less than 0.02 cfm at the highest sampling vacuum.

6.3 Instrumental Methods

Analyzer calibrations, system bias check and drift checks were completed before and after each sample run utilizing EPA Protocol 1 calibration gases.

The analyzer interference responses were determined in accordance with Section 5.4 of Method 20 and Section 7.2 through 7.6 of Method 6C.

The NOx analyzer NO_2 to NO converter efficiency is determined in accordance with Section 5.6 of Method 20.

7.0 Unit Operational Data

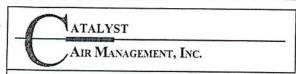
UGI will be responsible for the operation of the units and CEMS. UGI personnel will provide documentation for operational conditions maintained during the testing. The CEMS data acquisition and handling system (DAHS) will provide emissions data to determine the system relative accuracy.

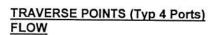
8.0 Test Report

The test report will be submitted that will provide the results of the test program and all the pertinent information pertaining to the individual tests. The report will include the following:

- Unit description
- Sampling location and test points
- Unit operating data and production data
- Test personnel
- Test results and discussion of the test program
- Raw test data
- Equipment calibration and test method QA
- Sample Calculations

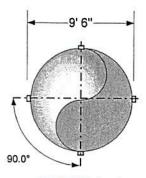
Figures





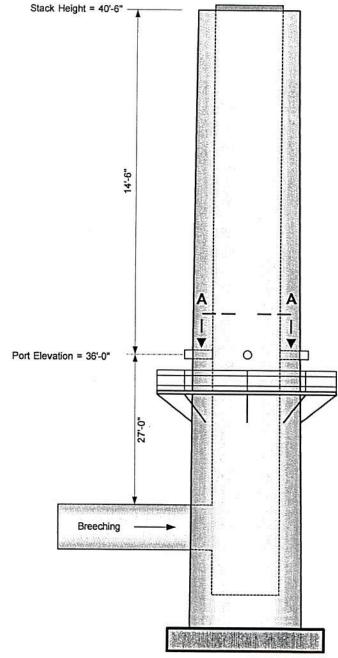
(Inches) from inside of stack.

- 1. <u>3.6"</u> 2. <u>12.0"</u>
- 3. <u>22.1"</u> 4. <u>36.8"</u>

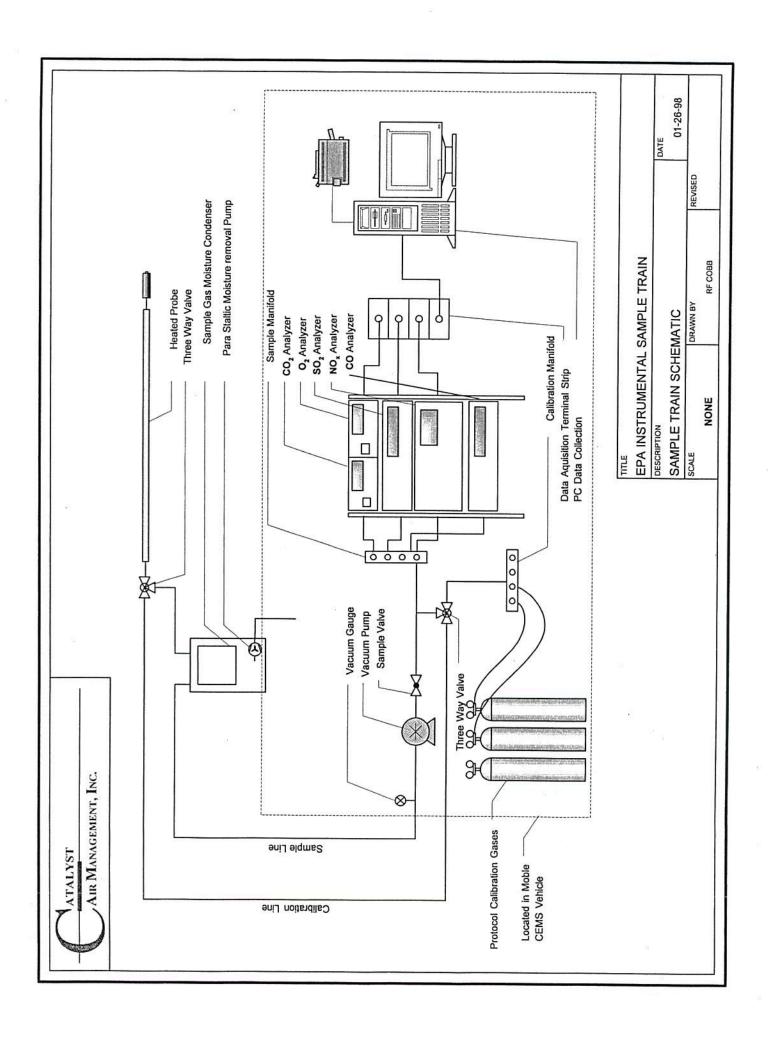


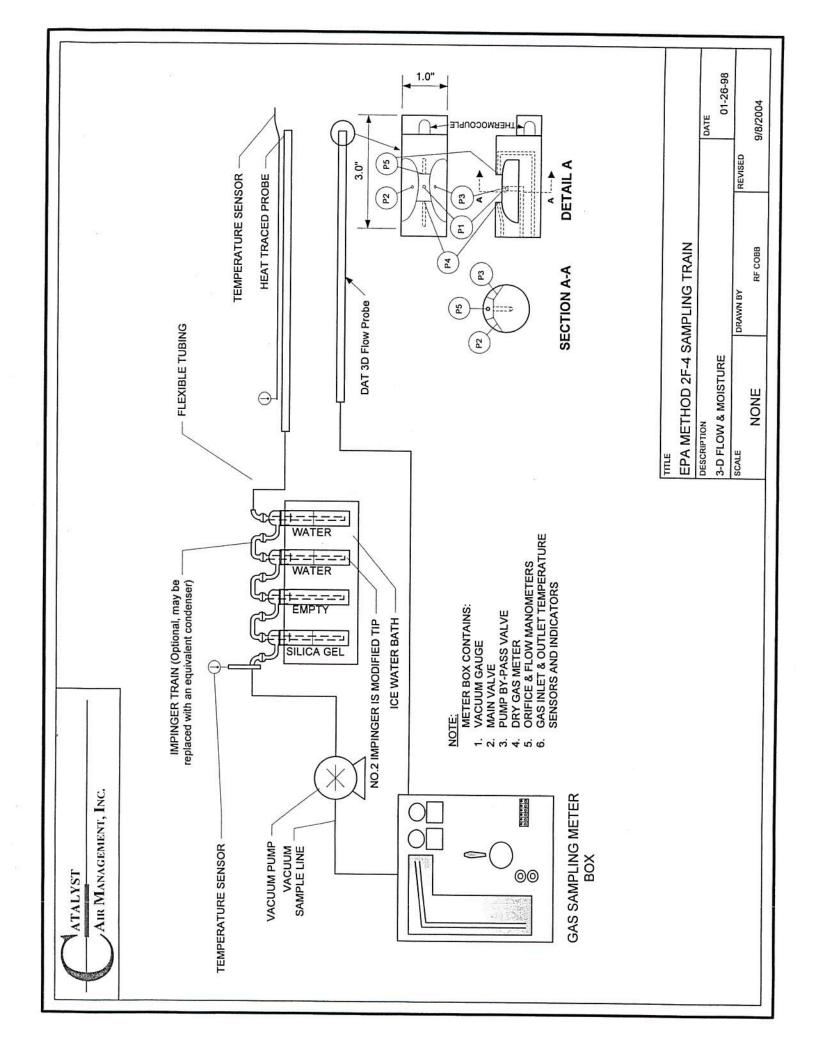
SECTION A - A

TRAVERSE POINTS (Typ 4 Ports) (meters) from inside of stack.



TITLE				
UGI DEVELOPME	NT COMPANY - HUNLO	CK STATION		
DESCRIPTION		DATE		
UNIT 6 TEST POR	T CONFIGURATION	7/24/		
SCALE	DRAWN BY	REVISED		
NONE	MJ TAYLOR			





Field Data Sheets

CATALYST AIR M TESTING SERVIC	IANAGE ES GRO	MENT Ir UP	ıc.		DATE:	pr		RUN#:		
CLIENT:					PLANT:			UNIT:		2
Analyzer:			O2	r:		CO2		1	NOx	ž.
Scale:		,						a.•		î D
Cylinder Value	TIME:	Zero Gas:	Mid-Range:	High-Range:	Zero Gas:	Mid-Range:	High-Range:	Zero Gas:	Mid-Range:	High-Range:
Analyzer Calib.										
System Calib. Run 1 Start: End:										
System Calib. Run 2 Start: End:			i i							
System Calib. Run 3 Start: End:										
System Calib. Run 4 Start: End:										
System Calib. Run 5 Start: End:										
System Calib. Run 6 Start: End:										
System Calib. Run 7 Start: End:		I								
System Calib. Run 8 Start: End:										-
System Calib. Run 9 Start: End:										
System Calib. Run 10 Start: End:										
System Calib. Run 11 Start: End:										
System Calib. Run 12 Start: End:										
Liiu.										

CATALYST AIR M	ANAGEMENT Inc
TESTING SERVICE	ES GROUP

PAGE 2

TESTING SERVICES GROUP				DATE:			RUN #:				
Analyzer:			SO ₂				2				
Scale:							71 6				15
			-	ii Taasi			±1	200			ř.
	TIME:	Zero Gas:	Mid-Range:	High-Range:	Zero Gas:	Low-Range:	Mid-Range:	High-Range:	Zero Gas:	Mid-Range:	High-Range:
Cylinder Value	[**	- "
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Analyzer Calib.	L					1					
System Calib.	Г					1					
Run 1 Start:											
End:											
System Calib.											
Run 2 Start:	====				Name of the last o						
End:					Y						
System Calib.											
Run 3 Start:			-		54						
End:											
System Calib.											
Run 4 Start: End:											
End: L System Calib.											
Run 5 Start:					<u> </u>						
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System Calib.						Г			ГТ	Т	
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Run 10 Start:					<u>.</u> —						
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Run 11 Start:											
End:											
System Calib.				1							
Run 12 Start:					1						
End:						855. (4. 77). 150					
	L										

CATALYST AIR MANAGEMENT Inc. TESTING SERVICES GROUP

FLOW DATA

FLOW DA	TA						
	CLIENT:			RUN #	PROBE ID:		
				LOAD	TC ID:		
				STATIC	Pre Leak:		
	UNIT: _		o B	BAROMETER	Post Leak:		
	03						
		TIME (24 HR.	SAMPLE	STACK TEMP	Yaw Angle	Velocity Pressure	Pitch
		CLOCK)	POINT	°F	degrees	(P1-P2)	Pressure (P4-P5)
	F	CEOCIE)	102.1		degrees	(1112)	(14-13)
							81
					i		
				X			
					la company of the com		
			•				
				Name of the last o			
MOISTUR	E DATA						
	Meter Box:			Y	i:	ž.	
	E+ 1598.0						
	POINT	CLOCK	RUN	DRY GAS	ORIFICE	METER TEMP	°F
	10	TIME	TIME	METER	PRESSURE	INLET	OUTLET
			0		1		
			5				
			10				
			15				
	_		20				
			25				
			30	9			
	IMPINGERS				SILICA GEL		
	IMPINGERS			- t ⊇3	FINAL WEIGHT		
	FINAL VOLUME INITIAL VOLUME			7 4	INITIAL WEIGHT NET WEIGHT		
				<u> </u>	HEI WEIGHT		
	NET VOLUM	Ξ		=3 =3	IMPINGERS		
					SILICA GEL		

TOTAL MOISTURE

Sample Calculations

SAMPLE EQUATIONS FOR CEMS RELATIVE ACCURACY TEST AUDITS

CALCULATIONS FOR FLUE GAS VOLUME AND MOISTURE

Time	Dry Gas Meter Ft ³	Pitot ΔP In. H ₂ O	Orifice ΔH In. H_2O	Dry Gas Temp.°F In Out	Flue Gas Static Pressure In. H ₂ O	Stack Temp °F
T	V_{m}	Δр	ΔΗ	TMI TMO	P_{g}	T_s

- 1. $P_{bar} = Barometric Pressure (in. Hg)$
- 2. TT = Net Sampling Time (minutes)
- 3. $V_m = V_m$ Final V_m Initial = Sample Gas Volume (Ft³)
- 4. T_m = Average Dry Gas Temperature at Meter (°F)

$$T_{m} = \frac{Avg.TMI + Avg.TMO}{2}$$

- 5. $\Delta p = \text{Velocity head of stack gas (in. H₂O)}$
- 6. $\Delta H = \text{Average Orifice Pressure Drop (in.H₂O)}$
- Volume of dry gas sampled at standard conditions^a (DSCF)

$$V_{m(std)} = \frac{(17.64)(V_m)(Y)\left(P_{bar} + \frac{\Delta H}{13.6}\right)}{(T_m + 460)}$$

- 8. V_{lc} = Total Water Collected = gm H_2O Silica gel + ml Imp. H_2O = ml
- Volume of water vapor at standard conditions^b (SCF)

$$V_{w(std)} = 0.0471(V_{lc}) = SCF$$

10. Percent moisture in flue gas

$$\% M = \frac{100(V_{w(std)})}{V_{m(std)} + V_{w(std)}}$$

11. Mole fraction of water vapor in flue gas

$$B_{ws} = \frac{\% M}{100}$$

12. Molecular Weight of dry flue gas

$$M_d = 0.44(\%CO_2) + 0.32(\%O_2) + 0.28(\%N_2 + \%CO)$$

13. Molecular weight of wet flue gas

$$M_s = M_d (1 - B_{ws}) + 18(B_{ws})$$

14. $A = Cross-sectional area of stack (Ft^2)$

$$\frac{\pi r^2}{144}$$

15. $P_s = \text{Flue gas pressure (in, Hg)}$

$$P_s = P_{bar} + P_g$$

NOTE:
$$P_g(Hg) = \frac{P_g(in. H_2 O)}{13.6}$$

16. $T_s = Absolute stack temperature (°R)$

$$T_s = 460 + t_s$$

17. Flue velocity at stack conditions (FT/SEC)

$$V_{s} = (K_{p})(C_{p}) \left[\left(\sqrt{\Delta p} \right) avg \right] \sqrt{\frac{T_{s}(avg)}{P_{s} * M_{s}}}$$

 C_p = pitot tube coefficient K_p = pitot tube constant = 85.49ft/sec 18. Flue gas volumetric flow rate at standard conditions^b (SCFM)

$$Q_s = (V_s)(A) \left(\frac{528}{T_s(avg.)}\right) \left(\frac{P_s}{29.92}\right) (60)$$

19. Flue gas volumetric flow rate at standard conditions^c (DSCFM)

$$Q_{sd} = (1 - B_{ws})(V_s)(A) \left(\frac{528}{T_s(avg.)}\right) \left(\frac{P_s}{29.92}\right) (60)$$

20. Flue gas volumetric flow rate at stack conditions (ACFM)

$$Q_a = (V_s)(A)(60)$$

NOTES:

^aDry standard cubic feet at 68°F, 29.92 in. Hg

^bStandard conditions at 68°F, 29.92 in. Hg

^cDry standard cubic feet per minute at 68°F, 29.92 in. Hg

CALCULATIONS FOR 3-D PITOT TUBE VELOCITY

RESULTANT ANGLE Ri

$$R_I = arc cosine [(cosine Yi) * (cosine Pi)]$$

AXIAL VELOCITY

$$V_I = Kp * F_2 * [Ti (P_1-P_2)/(Pa * Ms)]^{0.5} * (cosine Yi) * (cosine Pi)$$

V_i = Velocity at point i, fps Kp = Conversion factor 85.49

Ti = Absolute stack temperature at point i, °RP₁-P₂ = Pressure differential from 3-D probe, in H₂O

Pa = Absolute stack pressure, in Hg

Ms = Molecular weight of stack gas, lb/lb-mole F_2 = 3-D pitot coefficient, [(Pt-Ps)/ P_1 - P_2)]

Pt-Ps = Velocity pressure from standard type pitot, in H_2O

C_{ps} = Standard pitot tube coefficient

F-FACTOR DETERMINATION

THE WET F-FACTOR (Fw):

Includes all components of combustion

$$F_{_{w}} = \frac{10^{6} Btu / mmBtu [5.57(\% H) + 1.53(\% C) + 0.57(\% S) + 0.14(\% N) - 0.46(\% O_{_{2}}) + 0.21(\% H_{_{2}}O)]}{GCV_{_{wet}}}$$

THE DRY F-FACTOR (F_d):

Includes all components of combustion less water

$$F_{d} = \frac{10^{6} Btu / mmBtu \Big[3.64 \big(\% H\big) + 1.53 \big(\% C\big) + 0.57 \big(\% S\big) + 0.14 \big(\% N\big) - 0.46 \big(\% O_{2}\big) \Big]}{GCV_{dry}}$$

THE CARBON F-FACTOR (Fc):

Includes only Carbon Dioxide

$$F_c = \frac{10^6 Btu / mmBtu[0.321(\%C)]}{GCV_{dry}}$$

References for the above equations (i.e. %H, %C, %N, %S, %O₂) can be found in 40 CFR Part 60, Appendix A, Method 19.

CALCULATION FOR GAS CONCENTRATION

GAS CONCENTRATION (Cgas)

$$C_{gas} = \left(\overline{C} - C_0\right) \left(\frac{C_{ma}}{C_m - C_o}\right)$$

 C_{gas} = Effluent gas concentration, ppm

 \overline{C} = Average gas concentration indicated by gas analyzer, dry basis, ppm

C₀ = Average of initial and final system calibration bias check responses for the zero gas, ppm

C_m = Average of initial and final system calibration bias check responses for the upscale calibration gas, ppm

C_{ma} = Actual concentration of the upscale calibration gas, ppm

GAS CONCENTRATION @ 15% O2 (Cgas @ 15% O2)

$$C_{gas}$$
 @ 15% $O_2 = C_{gas} * ((20.9-15)/(20.9-\%O_2))$

GAS CONCENTRATION @ 7% O2 (Cgas @ 7% O2)

$$C_{gas} @ 7\% O_2 = C_{gas} * ((20.9-7)/(20.9-\%O_2))$$

GAS CONCENTRATION @ 3% O_2 (C_{gas} @ 3% O_2)

$$C_{gas} @ 3\% O_2 = C_{gas} * ((20.9-3)/(20.9-\%O_2))$$

LBS/MMBTU CALCULATIONS USING THE F-FACTOR

1. EMISSION RATE E(lb/mmbtu), O2 based

$$E(lb/mmbtu) = C \times F_d \left(\frac{20.9}{20.9 - \%O_2} \right)$$

Where:

C(lb/dscf) = Pollutant concentration (ppm) x conversion factor.

Conversion Factors:

$$NOx = 1.194 \times 10^{-7}$$

$$SO_2 = 1.660 \times 10^{-7}$$

$$CO = 7.274 \times 10^{-8}$$

$$C_3H_8 = 1.145 \times 10^{-7}$$

 $F_d(dscf/mmbtu) = "F"$ Factor for fuel type, (Ref. EPA Method 19)

$$F_d$$
 (Coal) = 9780

$$F_d$$
 (Gas) = 8710

$$F_d$$
 (Oil) = 9190

2. EMISSION RATE E(lb/mmbtu), CO2 based

$$E(lb/mmbtu) = C \times F_c \left(\frac{100}{\%CO_2}\right)$$

Where:

C(lb/dscf) = Pollutant concentration (ppm) x conversion factor.

Conversion Factors:

$$NOx = 1.194 \times 10^{-7}$$

$$SO2 = 1.660 \times 10^{-7}$$

$$CO = 7.274 \times 10^{-8}$$

$$C_3H_8 = 1.145 \times 10^{-7}$$

 $F_c(dscf/mmbtu) =$ "F" Factor for fuel type, (Ref. EPA Method 19)

$$F_d$$
 (Coal) = 1800

$$F_d(Gas) = 1040$$

$$F_d$$
 (Oil) = 1420

CALCULATION OF RELATIVE ACCURACY

ARITHMETIC MEAN (OF THE DIFFERENCE, {d}, OF A DATA SET)

$$\overline{d} = \frac{1}{n} \sum_{i=1}^{n} d_i$$

Where n = Number of data points.

ALGEBRAIC SUM (OF THE INDIVIDUAL DIFFERENCES, {d_i})

$$\sum_{i=1}^{n} d_{i}$$

STANDARD DEVIATION, S_d

$$S_{d} = \sqrt{\frac{\sum_{i=1}^{n} d_{i}^{2} - \left(\frac{\left(\sum_{i=1}^{n} d_{i}\right)^{2}}{n}\right)}{n-1}}$$

CONFIDENCE COEFFICIENT, CC

$$CC = t_{0.975} \frac{S_d}{\sqrt{n}}$$

For 9 tests $t_{0.975} = 2.306$

For 10 tests $t_{0.975} = 2.262$

For 11 tests $t_{0.975} = 2.228$

For 12 tests $t_{0.975} = 2.201$

RELATIVE ACCURACY, RA

$$RA = \frac{\left|\overline{d}\right| + \left|CC\right|}{\overline{RM}}x100$$